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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(21) International Application Number: PCT/US97/11296 (22) International Filing Date: 26 June 1997 (26.06.97) (30) Priority Data: 60/022,518 28 June 1996 (28.06.96) US (71) Applicant (for all designated States except US): E.I. DU PONT DE NEMOURS AND COMPANY [US/US]; 1007 Market Street, Wilmington, DE 19898 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): CICHA, Walter, Vladimir [CA/US]; Apartment 4-C, 502 Red Fox Lane, Newark, DE 19711 (US). MANZER, Leo, Ernest [US/US]; 714 Burnley Road, Wilmington, DE 19803 (US). (74) Agent: HEISER, David, E.; E.I. du Pont de Nemours and Company, Legal Patent Records Center, 1007 Market Street, Wilmington, DE 19898 (US).		(81) Designated States: BR, CN, IL, JP, RU, US, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: PHOSGENE MANUFACTURING PROCESS (57) Abstract A process for producing phosgene is disclosed which involves contacting a mixture comprising CO and Cl ₂ (e.g., at about 300 °C or less) with a silicon carbide catalyst having a surface area of at least 10 m ² .g ⁻¹ .		

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TITLE
PHOSGENE MANUFACTURING PROCESS
FIELD OF THE INVENTION

5 This invention relates to a process for the manufacture of phosgene by the reaction of chlorine (Cl_2) with carbon monoxide (CO) in the presence of a catalyst. More particularly, this invention relates to a process for the manufacture of phosgene with minimal production of the hazardous chemical, carbon tetrachloride.

BACKGROUND

10 The production of phosgene by the reaction of chlorine with carbon monoxide in the presence of a carbon catalyst is a well known process. The phosgene produced by this process will typically contain 400 to 500 ppm by weight carbon tetrachloride. This amount, needs to be evaluated on the basis of the total world-wide production of phosgene which is about ten billion pounds
15 (4.5×10^9 kg), corresponds to co-production of about 4 to 5 million pounds (1.8×10^6 kg to 2.3×10^6 kg) of carbon tetrachloride with the phosgene.

A Japanese patent publication (Kokoku) Patent No. Hei 6[1994]-29129 discloses that the amount of carbon tetrachloride produced during the phosgene manufacturing process can be reduced by about 50% to about 150 ppm by weight
20 by using an activated carbon which has been washed with an acid and which contains a total of 1.5 wt.% or less of metal components comprised of transition metals, boron, aluminum and silicon.

Carbon tetrachloride has been shown to have both significant ozone depletion and global warming potentials. Therefore, there is an interest in
25 developing phosgene processes in which the amount of carbon tetrachloride impurity is minimized.

Silicon carbide has long been known as a material which has high thermal and chemical stability, has excellent heat and electrical conducting properties and as an abrasive which is almost as hard as diamond. Silicon carbide can be
30 commercially prepared electrochemically using the Acheson process. The product so produced has a surface area of less than $1 \text{ m}^2/\text{g}$; its use as a catalyst support has been limited partly because of this low surface area. Recently, high surface area (60 to $400 \text{ m}^2/\text{g}$) silicon carbides have been prepared (M. J. Ledoux et al., J. Catal., 114, 176-185 (1988)). These high surface area materials are used as
35 catalyst supports or carriers.

SUMMARY OF THE INVENTION

A process for producing phosgene is provided which comprises contacting a mixture comprising carbon monoxide and chlorine at a temperature of about

300°C. or less. with a catalyst comprising silicon carbide and having a surface area of at least $10 \text{ m}^2 \cdot \text{g}^{-1}$.

DETAILED DESCRIPTION

The present invention relates to improving the production of phosgene produced by contacting carbon monoxide and chlorine. Surprisingly, we have found that silicon carbide itself can be used as a catalyst for the manufacture of phosgene. The improvement can be employed in connection with the operating conditions used for any of the carbon-based processes previously used commercially or described in the art (e.g., those processes disclosed in U.S. Patent Nos. 4,231,959 and 4,764,308 for the production of phosgene).

Phosgene is commercially manufactured by passing carbon monoxide and chlorine over activated carbon. The reaction is strongly exothermic and is usually done in multitubular reactors to more effectively control the reaction temperature. Carbon monoxide is added in at least a stoichiometric amount (often in stoichiometric excess) to minimize the free chlorine content of the product phosgene.

The reaction temperature and the silicon carbide are chosen to provide phosgene which contains about 300 ppm or less by weight of carbon tetrachloride. Preferably, the phosgene contains less than about 250 ppm or less by weight of carbon tetrachloride; most preferably, the phosgene contains less than about 100 ppm or less by weight of carbon tetrachloride.

Any silicon carbide-containing catalyst with a surface area greater than about $10 \text{ m}^2/\text{g}$ (e.g., about $20 \text{ m}^2/\text{g}$, or more) may be used in the process of this invention. However, silicon carbide compositions having surface areas greater than about $100 \text{ m}^2/\text{g}$, prepared by the methods disclosed in U.S. Patent No. 4,914,070 (incorporated herein by reference) are especially preferred. A silicon content of at least about 5 weight % is preferred. More preferably the silicon content is at least about 10 weight %. Of note are embodiments where the catalyst is manufactured using a process which comprises contacting silicon monoxide with finely divided carbon (see e.g., U.S. Patent No. 4,914,070). Use of a carbon which has an ash content of less than about 0.1 wt. % to produce the silicon carbide is preferred.

A preferred silicon carbide catalyst is prepared by a process comprising reacting vapors of silicon monoxide, SiO, on carbon by the steps of:

- (a) generating vapors of SiO in a first reaction zone by heating a mixture of SiO_2 and Si at a temperature of between 1100°C and 1400°C, under a pressure of between 0.1 and 1.5 hPa; and
- (b) contacting in a second reaction zone at a temperature of between 1100°C and 1400°C, the SiO vapors generated in said first reaction zone with finely divided reactive carbon with a specific surface area that

is equal to or greater than $200 \text{ m}^2 \cdot \text{g}^{-1}$. Examples of suitable reactive carbons include graphite pellets obtained by powder agglomeration; and activated carbon such as powdered activated carbon obtained by crushing granules of activated carbon.

- 5 The silicon carbide surface area as determined by BET measurement is preferably greater than about $100 \text{ m}^2/\text{g}$ and more preferably greater than about $300 \text{ m}^2/\text{g}$.

It is known from dissociation equilibria that at 100°C , phosgene contains about 50 ppm chlorine; and that at 200°C , about 0.4%, at 300°C , about 5% and at
10 400°C about 20% of the phosgene is dissociated into carbon monoxide and chlorine. Also, the higher the reaction temperature the more carbon tetrachloride is produced. Accordingly, the temperature of the reaction is generally about 300°C or less (e.g., in the range of from 40°C to 300°C). Preferably, the temperature of the process is from about 50°C to 200°C ; more preferably from
15 about 50°C to 150°C . The phosgene produced by the process of this invention typically contains about 300 ppm by weight or less of carbon tetrachloride, based upon phosgene (i.e., 300 parts by weight CCl_4 per million parts by weight COCl_2 , or less) even at a temperature of 300°C . Preferably, the reaction temperature and the silicon carbide are chosen to provide phosgene which contains less than about
20 250 ppm by weight of carbon tetrachloride; and more preferably, are chosen to provide phosgene which contains less than 100 ppm by weight of carbon tetrachloride, based upon phosgene. Of note are embodiments where the reaction time and temperature are controlled to provide a carbon tetrachloride concentration of 100 ppm or less based upon the total product stream.

- 25 Without further elaboration, it is believed that one skilled in the art can, using the description herein, utilize the present invention to its fullest extent. The following preferred specific embodiments are, therefore, to be construed as merely illustrative, and does not constrain the remainder of the disclosure in any way whatsoever.

30

EXAMPLES

The following procedures were used in Example 1 and Comparative Example A for catalyst testing and for product analysis.

General Catalyst Testing Procedure

- 35 A $1/2"$ (1.27 mm) O.D. x 15" (381 mm) Inconel® 600 nickel alloy tube containing a 100 mesh (0.015 mm) Monel® nickel alloy screen was used as the reactor. The reactor was charged with about 2.5 mL to about 8 mL of silicon carbide catalyst and heated to 300°C . This was the temperature used for all the examples.

A 1:1 molar ratio mixture of carbon monoxide and chlorine was passed over the catalyst. The contact times were between 0.9 to 12 seconds. The experimental results are shown in Tables 1 and A.

General Analytical Procedure

- 5 The reactor effluent was sampled on-line with a Hewlett Packard HP 5890 gas chromatograph using a 105 m long, 0.25 mm I.D. column containing Restak™ RTX-1 Crossbond 100% dimethyl polysiloxane. Gas chromatographic conditions were 50°C for 10 minutes followed by temperature programming to 200°C at a rate of 15°C/minute. The smallest amount of carbon tetrachloride that could be
- 10 quantitatively identified was about 40 ppm by weight for Example 1 and about 80 ppm by weight for Comparative Example A. Testing results are shown in Tables 1 and A.

TABLE 1

Example	CCl ₄ Conc. ¹ ppm	COCl ₂ ² mole %	Si Content wt. %	Surface Area m ² /g
1	50	>77	29.0	480

¹By weight as ppm of the product stream. The value shown is an average taken over 7 hours and is a high-end estimate

²Fraction in product gas stream over initial 7 hours of the run. The contact time was 0.9 to 1.3 seconds

TABLE A

Comparative Example	CCl ₄ Conc. ¹ ppm	COCl ₂ ² mole %	Si Content wt. %	Surface Area m ² /g
A	<80	4.0	70.0 ³	0.062

¹By weight as ppm of the product stream. The value shown is an average taken over 7 hours and is a high-end estimate

²Fraction in product gas stream over initial 7 hours of the run. The contact time was 8 to 12 seconds

³Calculated based on stoichiometry

What is claimed is:

1. A process for producing phosgene, comprising:
contacting a mixture comprising CO and Cl₂ at a temperature of
about 300°C, or less, with a catalyst comprising silicon carbide and having a
5 surface area of at least 10 m²•g⁻¹.
2. The process of Claim 1 wherein the catalyst is manufactured using a
process which comprises contacting silicon monoxide with finely divided carbon.
3. The process of Claim 2 wherein the carbon used to produce the silicon
carbide has an ash content of less than about 0.1 wt %.
- 10 4. The process of Claim 1 where the silicon carbide is obtained by
(a) generating vapors of SiO in a first reaction zone by heating a mixture of SiO₂
and Si at a temperature of between 1100°C and 1400°C, under a pressure of
between 0.1 and 1.5 hPa; and (b) contacting in a second reaction zone at a
temperature of between 1100°C and 1400°C, the SiO vapors generated in said first
15 reaction zone with finely divided reactive carbon with a specific surface area that
is equal to or greater than 200 m²•g⁻¹.

INTERNATIONAL SEARCH REPORT

Inter. Application No

PCT/US 97/11296

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C01B31/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 014, no. 131 (C-0700), 13 March 1990 & JP 02 006307 A (IDEMITSU PETROCHEM CO LTD), 10 January 1990, see abstract ---	1
A,P	DATABASE WPI Section Ch, Week 9719 Derwent Publications Ltd., London, GB; Class A23, AN 97-209105 XP002044695 & JP 09 059 012 A (MITSUBISHI GAS CHEM CO INC), 4 March 1997 see abstract --- -/-	1

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	EP 0 003 530 A (STAUFFER CHEMICAL CO) 22 August 1979 see claim 1 & US 4 231 959 A cited in the application -----	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/11296

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